

Rifampin Level (PKRIF)

Overview

Method Name

HPLC

NY State Available

No

Specimen

Specimen Type Serum

# **Specimen Required**

Container/Tube: Red Top

Preferred: Red top tube

Acceptable: Serum gel tube

Specimen Volume: 2 mL

**Collection Instructions**: Draw blood in a plain, red-top tube(s). Separate serum from cells immediately by centrifugation and aliquot into a polypropylene or similar plastic tube. Send 2 mL of serum frozen in plastic vial. Note:

1. The following information is required:

A. Specimen Type (source)

B. Dose (specify PO, IV, IM)

C. Date and time of last dose (for IV start/end time)

2. If the time of last dose and the blood draw are not accurately recorded, accurate interpretation of the concentration is not possible.

# Specimen Minimum Volume

0.5 mL

# **Reject Due To**

Hemolysis	Mild OK; Gross reject
Lipemia	NA
Icterus	NA
Other	Samples thawed greater than 6 hours.

# Specimen Stability Information

**Test Definition: FRIFA** 



Specimen Type	Temperature	Time	Special Container
Serum	Frozen	365 days	

## Clinical & Interpretive

ABORATORIES

## **Reference Values**

mcg/mL

#### Interpretation

The target range for mycobacterial infections is 8 to 24 mcg/mL 2 hours after oral dose or 2 hours after the end of intravenous infusion. Samples drawn later than 2 hours after the dose will often display concentrations below the stated range.

Rifampin generally should be given as a single daily dose. If the patient is receiving 2 small daily doses, consider combining the doses and rechecking the concentration.

Rifampin absorption may be reduced by food. Take on an empty stomach if possible.

Rifampin does not have clear concentration-related toxicity and most patients tolerate concentrations above the stated range without difficulty.

Hepatic dysfunction may produce elevated rifampin concentrations. Rifampin concentrations greater than 50% above the range may warrant a dose reduction of 150 to 300 mg.

If the time of the dose and the blood draw were not accurately recorded, accurate interpretation of the concentration is not possible.

## Performance

PDF Report No

Day(s) Performed Monday through Friday

Report Available 10 to 14 days

Performing Laboratory Location

National Jewish Health



# **Test Definition: FRIFA**

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# Fees & Codes

## Fees

- Authorized users can sign in to <u>Test Prices</u> for detailed fee information.
- Clients without access to Test Prices can contact <u>Customer Service</u> 24 hours a day, seven days a week.
- Prospective clients should contact their account representative. For assistance, contact Customer Service.

#### **Test Classification**

The performance characteristics for this test have been validated by Advanced Diagnostic Laboratories at National Jewish Health. It has not been cleared or approved by the US Food and Drug Administration. The results are not intended to be used as the sole means for clinical diagnosis or patient management decisions. This laboratory is certified under the Clinical Laboratory Improvement Amendments of 1988 (CLIA 88) as qualified to perform high complexity clinical laboratory testing.

#### **CPT Code Information**

80299

## LOINC<sup>®</sup> Information

Test ID	Test Order Name	Order LOINC <sup>®</sup> Value
FRIFA	Rifampin Level	Not Provided

Result ID	Test Result Name	Result LOINC <sup>®</sup> Value
Z5803	Dose	Not Provided
Z5804	Date and Time of Last Dose	Not Provided
Z5778	Rifampin Level	4021-2
Z5840	Comment:	Not Provided
Z5865	Specimen Type:	Not Provided